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S Ma E	r ຸ U.S. Patent and	PTO/SB/21 (08-03) Approved for use through 08/30/2003. OMB 0651-0031 Trademark Office; U.S. DEPARTMENT OF COMMERCE
TRANSMITTAL FORM (to be used for all correspondence after initial for	Application Number Filing Date First Named Inventor Art Unit Examiner Name	nformation unless it disolars a valid OMB control number. 10/010, 065 December 5, 2001 Ceith D. ALLEN 1632 Dertoglio, Valerie E R-648
Fee Transmittal Form Fee Attached Amendment/Reply After Final	ENCLOSURES (Check all that app Drawing(s) Licensing-related Papers Petition Petition to Convert to a Provisional Application Power of Attorney, Revocation	
Extension of Time Request Express Abandonment Request Information Disclosure Statement Certified Copy of Priority Document(s) Response to Missing Parts/	Change of Correspondence Address Terminal Disclaimer Request for Refund CD, Number of CD(s) Remarks	Other Enclosure(s) (please Identify below): RECEIVED DEC 1 9 2003
Firm or Individual name Kelly L. Que		OR AGENT
I hereby certify that this correspondence is be sufficient postage as first class mail in an env	RTIFICATE OF TRANSMISSION/MA ing facsimile transmitted to the USPTO or depr	AILING posited with the United States Postal Service with Property Propert
Typed or printed name Kelly L Signature Kelly J This collection of information is required by 37 CFR	**************************************	enefit by the public which is to file (and by the USPTO to

process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to 12 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450. DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

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FEE TRANSMITTA			Application Number			1 / -	
for EV 2004			Filing Date			December 5, 2001	
for FY 2004			First Named Inventor		Invent	for Keith D. ALLEN	
Effective 10/01/2003. Patent fees are subject to annual revision.			Examiner Name		ame	Bertoglio, Valerie	; E 1
Applicant claims small entity status. See 37 CFR 1.27			Art Unit			1632	
TOTAL AMOUNT OF PAYMENT	(\$) 55.00		Attorr	ney Do	cket No	1 110	0~
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The Director is authorized to: (check all that apply) Charge fee(s) indicated below Credit any overpayments		1812	2,520			or filing a request for ex parte reexamination	1
Charge any additional fee(s) or any underpayment of fee(s)		1804	920*	1804	920* F	Requesting publication of SIR prior to Examiner action	
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to the above-identified deposit account.			,		, I	Examiner action	55.00
FEE CALCULATION		1251	-	2251		Extension for reply within first month	33.55
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SUBTOTAL (1) (\$)	1452	110	2452	55 1	Petition to revive - unavoidable	
2. EXTRA CLAIM FEES FOR UTILITY AND REISSUE			1,330	2453		Petition to revive - unintentional	—
Extra Claims	Fee from _	1501 1502	1,330 480	2501 2502		Utility issue fee (or reissue) Design issue fee	
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1202 18 2202 9 Claims in ex		1809	770	2809	385	Filing a submission after final rejection	
	t claims in excess of 3 pendent claim, if not paid	1910	770	2015		(37 CFR 1.129(a)) For each additional invention to be	
<u>'</u>	independent claims	1810	770	2810		examined (37 CFR 1.129(b))	
over origin	nal patent	1801		2801		. , ,	
	claims in excess of 20 original patent	1802	900	1802		Request for expedited examination of a design application	

**or number previously paid, if greater, For Reissues, see above (Complete (if applicable)) SUBMITTED BY Registration No. L. Quast (650) 569-5100 52,141 Name (Print/Type) Kelly Telephone (Attorney/Agent) 12/10/03 Kelly fluart Date Signature

Other fee (specify)

*Reduced by Basic Filing Fee Paid

SUBTOTAL (3)

55,00

WARNING: Information on this form may become public. Credit card information should not be included n this form. Provide credit card information and authorization on PTO-2038.

This collection of information is required by 37 CFR 1.17 and 1.27. The information is required to obtain or retain a benefit by the public which is to file (and by the USPTO to process) an application. Confidentiality is governed by 35 U.S.C. 122 and 37 CFR 1.14. This collection is estimated to take 12 minutes to complete, including gathering, preparing, and submitting the completed application form to the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete this form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Trademark Office, U.S. Department of Commerce, P.O. Box 1450, Alexandria, VA 22313-1450, DO NOT SEND FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450.

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SUBTOTAL (2)



UNITED STATES PATENT AND TRADEMARK OFFICE

UNITED STATES DEPARTMENT OF COMMERCE UNITED STATES DEPARTMENT OF COMMIT United States Patent and Trademark Office Address COMMISSIONER FOR PATENTS P.O. Box 1450 Absundita, Vignia 22313-1450 www.mpto.gov

ATTORNEY DOCKET NO.

R-648

ART UNIT

APPLICATION NO. FILING DATE FIRST NAMED INVENTOR 10/010,065 12/05/2001 Keith D. Allen 7590 08/13/2003 DELTAGEN, INC. 740 Bay Road Redwood City, CA 94063

EXAMINER BERTOGLIO, VALARIE E

CONFIRMATION NO.

2751

PAPER NUMBER

1632 DATE MAILED: 08/13/2003

Please find below and/or attached an Office communication concerning this application or proceeding.

IPE	<u> </u>	Application No.	Applicant(s)			
		10/010,065	ALLEN ET AL.			
1 5 2003	Office Action Summary	Examiner	Art Unit			
a	<i>E</i>	Valarie Bertoglio	1632			
Pariod f	- Th MAILING DATE of this communication app r Reply	pears on the cover sheet	with the correspondence address -			
A SH THE - Exte afte - If th - If No - Faili - Any	HORTENED STATUTORY PERIOD FOR REPLIMAILING DATE OF THIS COMMUNICATION. The ensions of time may be available under the provisions of 37 CFR 1.1 and the state of this communication. The period for reply specified above is less than thirty (30) days, a replimate of the period for reply is specified above, the maximum statutory period to the period for reply within the set or extended period for reply will, by statute reply received by the Office later than three months after the mailing the patent term adjustment. See 37 CFR 1.704(b).	36(a). In no event, however, may y within the statutory minimum of t will apply and will expire SIX (6) Min. cause the application to become	a reply be timely filed hirty (30) days will be considered timely. ONTHS from the mailing date of this communication. ABANDONED (35 U.S.C. § 133).			
Status _						
1)[Responsive to communication(s) filed on 14 A	A <i>pril 2003</i> .				
2a)⊠	, 	is action is non-final.				
3) <u> </u>	Since this application is in condition for allows closed in accordance with the practice under ion of Claims					
4)	Claim(s) <u>1-4,11-17 and 35-71</u> is/are pending in	n the application.				
	4a) Of the above claim(s) <u>1-4,11-17 and 35-56</u>	is/are withdrawn from co	onsideration.			
5)	Claim(s) is/are allowed.					
6)⊠	Claim(s) 57-71 is/are rejected.					
7) Claim(s) is/are objected to.						
	Claim(s) are subject to restriction and/or	election requirement.	·			
	ion Papers					
	The specification is objected to by the Examine					
10)	The drawing(s) filed on is/are: a)⊠ accep	ted or b) objected to by	the Examiner.			
	Applicant may not request that any objection to the	- , .				
11) 🔲	The proposed drawing correction filed on		disapproved by the Examiner.			
, -	If approved, corrected drawings are required in rep					
	The oath or declaration is objected to by the Exa	aminer.				
	ınder 35 U.S.C. §§ 119 and 120					
13) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).						
a) All b) Some * c) None of:						
1. Certified copies of the priority documents have been received.						
2. Certified copies of the priority documents have been received in Application No						
* S	3. Copies of the certified copies of the priori application from the International Bur see the attached detailed Office action for a list of	eau (PCT Rule 17.2(a)).				
14)⊠ A	cknowledgment is made of a claim for domestic	priority under 35 U.S.C	. § 119(e) (to a provisional application)			
a	☐ The translation of the foreign language province the comment is made of a claim for domestic	visional application has t	peen received.			
ttachment						
) 🔲 Notice	e of References Cited (PTO-892) e of Draftsperson's Patent Drawing Review (PTO-948) nation Disclosure Statement(s) (PTO-1449) Paper No(s)	5) Notice of	Summary (PTO-413) Paper No(s). <u>12</u> . Informal Patent Application (PTO-152)			

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Response to Amendment

Applicant's arguments filed 04/14/2003, paper number 12, have been fully considered. Claims 5-10 and 18-34 have been cancelled. Claims 57-71 have been added. Claims 1-4,11-17 and 35-56 have been withdrawn. Claims 1-4,11-17, 35-71 are pending and claims 57-71 are under consideration in the instant action.

Specification

Applicants arguments to the objection to the specification on the grounds that the specification is unclear as to the phenotype of the claimed mice with regard to fasting blood glucose levels has been considered and was not found persuasive. The statement on page 62, lines 28-30 of the specification, disclosing that heterozygous mutant mice display decreased fasting glucose levels is contradictory to the statement on page 59, line 12 disclosing that heterozygous mutant mice have increased fasting blood glucose levels. Figure 4 supports the statement that heterozygous mutant mice had increased blood glucose levels. Applicants respond that Table 2 does not refer to fasting glucose levels but non-fasting glucose levels. However, the relevance of this argument is unclear. Examiner's explanation of the lack of clarity of the specification with regard to the phenotype of the mouse did not rely on Table 2. Applicants' response also gives evidence that heterozygous mice display reduced serum non-fasting glucose levels (page 4, lines 24-29), however, they fail to address the fact that the specification states that "heterozygous mutant mice had increased fasting blood glucose levels." (page 59, line 12-page 60, line 1). With evidence in the specification to support contradictory phenotypes, it cannot be determined which phenotype(s) the mice actually displayed.

Claim Rejections - 35 USC § 112, 1st paragraph

The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the

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art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

Claims 57-71 as newly added are rejected under 35 U.S.C 112, 1st paragraph as containing subject matter which was not described in the specification in such a way as to reasonably enable one skilled in the art to which it pertains, or with which it is most nearly connected, to make and/or use the invention. Claims 5-10 and 18-34 have been cancelled and thus, the rejection as it pertains to claims 5-10 and 18-34 is withdrawn. The previous rejection is, however, applicable to claims 57-71 for the reasons of record advanced on pages 7 and 8 of the previous office action mailed 01/07/2003. For clarity, the rejection is reiterated below.

1) The specification fails to enable knocking out any glucagon receptor gene other than that set forth by SEQ ID NO:1 (see page 8, 2nd paragraph of prior office action mailed 01/07/2003).

By use of the phrase "an endogenous glucagon receptor gene", claims 57 and 70, are drawn to any glucagon receptor gene, including orthologues of the mouse glucagon receptor set forth by SEQ ID NO:1. The specification only teaches one mouse glucagon receptor gene (SEQ ID NO: 1; page 10, lines 6-8). The specification does not provide adequate guidance for determining other mouse glucagon receptor genes or that other glucagon receptor genes exist in mouse or have the same function as the glucagon receptor gene disclosed. Therefore, a knockout of any glucagon receptor gene other than the glucagon receptor described in the specification would have different phenotypic effects that are not predictable (see pages 5-6 of the prior office action). Deleting the word "an" preceding the term "endogenous" in claims 57 and 70 would overcome this rejection.

2) The specification fails to enable making mice comprising a homozygous and/or heterozygous disruption of the glucagon receptor gene wherein the mice exhibit any phenotype

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(see page 5, paragraph 2; page 7, paragraph2; page 8, paragraph 1 of prior office action mailed 01/07/2003).

Claim 57 encompasses mice heterozygous for a disruption in the glucagon receptor gene wherein the mice have any phenotype. The phrase, "A transgenic mouse whose genome comprises a disruption in an endogenous glucagon receptor gene" encompasses both homozygous and heterozygous mice. The following phrase of the claim, "wherein where the disruption is homozygous, the transgenic mouse exhibits, relative to a wild-type mouse, a metabolic abnormality or a pancreatic abnormality" limits the recited phenotype to homozygous mice only. Therefore, the heterozygous mice encompasses by the claim have no recited phenotype and therefore, encompasses mice heterozygous for a disruption in the glucagon receptor gene wherein the mouse has any phenotype.

The specification teaches that mice heterozygous for a disruption in the glucagon receptor gene display both increased fasting blood glucose levels (sentence bridging pages 59-60) and decreased fasting glucose levels (page 62, lines 28-30), increased fasting insulin levels (Figure 11), reduced non-fasting serum glucose levels (page 59, line 2; Table 2) and mild to moderate hyperplasia and hypertrophy of the islet cells (page 56, lines 20-21). Therefore, the specification is enabling for heterozygous mice having only those phenotypes as described by the specification. As set forth in the art and described in said prior office action (page 5, paragraph 2), the phenotype of a transgenic animal was unpredictable at the time of filing. The specification does not overcome the unpredictability inherent in generating knockout mice such that <u>any</u> phenotype could be obtained and its use determined prior to learning the phenotype. The specification does not teach any phenotype for the claimed heterozygous mice other than those listed above. Without reciting a phenotype in claim 57 for the heterozygous mice, the claim encompasses heterozygous mice with other phenotypes not supported by the

specification. Without guidance as to how to obtain and use the heterozygous mice, it would require one of skill in the art at the time the invention was made, undue experimentation to determine how to obtain any phenotype in a mouse heterozygous for a disruption in the glucagon receptor gene or to use said mouse wherein the mouse has any phenotype.

Claim 70 encompasses a transgenic mouse whose genome comprises a homozygous disruption in an endogenous glucagon receptor gene wherein the mouse has any phenotype. There is no phenotype recited for the claimed mouse. The phenotype of reduced fertility recited in line 5 of the claim is the phenotype of the offspring of the claimed mouse and does not reflect any observable or detectable phenotype of the <u>claimed</u> mouse that correlates to the disruption in the glucagon receptor gene. The specification fails to overcome the unpredictability in the art as set forth above and in the prior office action. Accordingly, it would require one of skill in the art at the time the invention was made, undue experimentation to determine how to make and use the claimed mouse having any phenotype.

Claims 58-69 depend from parent claim 57 and also encompass both the homozygous heterozygous mice encompassed by the parent claim. Accordingly, the above rejection also applies to claims 58-69.

Claim 71 depends from parent claim 70 and also encompasses both the homozygous heterozygous mice encompassed by the parent claim. Accordingly, the above rejection also applies to claim 71.

Applicants argue that the rejections reiterated above are overcome by limitations introduced to newly added claims 57-71. This argument is not persuasive. Applicants' do not address how newly added claims 57 and 70 overcome the rejection to the broad genera of glucagon receptor genes encompassed by the phrase "an endogenous glucagon receptor".

Applicants' also argue that the specification is fully enabling for the newly added claims as they

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are limited to specific phenotypes. However, claim 57 is written such that it encompasses a heterozygous mouse exhibiting any phenotype. Claims 58-69 and 71 are dependent from claim 57 and encompass both heterozygous and homozygous mice wherein the mice have specific phenotypes. However, the phenotypes cited by claims 59-63, 65-69 are not supported by the specification. Furthermore, the homozygous mouse of claim 70 does have a phenotype. Therefore, applicants' arguments with respect to claims 57-71 are not persuasive the rejections reiterated above are maintained for the reasons of record.

Conclusion

Applicant's amendment necessitated the new ground(s) of rejection presented in this Office action. Accordingly, **THIS ACTION IS MADE FINAL**. See MPEP § 706.07(a). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Valarie Bertoglio whose telephone number is 703-305-5469. The examiner can normally be reached on Mon-Weds 6:00-2:30.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Deborah Reynolds can be reached on 703-305-4051. The fax phone numbers for

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the organization where this application or proceeding is assigned are 703-872-9306 for regular communications and 703-872-9307 for After Final communications.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is 703-308-1234.

PETER PARAS

Valarie Bertoglio Examiner Art Unit 1632 Page 7

Application No. Applicant(s) 10/010,065 ALLEN ET AL. Intervi w Summary Examiner Art Unit Valarie Bertoglio 1632 All participants (applicant, applicant's representative, PTO personnel): (1) Valarie Bertoglio. (2) Nicole Verona. Date of Interview: 02 June 2003. Type: a) ✓ Telephonic b) ✓ Video Conference c) Personal [copy given to: 1) applicant 2) applicant's representative Exhibit shown or demonstration conducted: d) Yes e) ∏ No. If Yes, brief description: _____. Claim(s) discussed: 57-85. Identification of prior art discussed: Agreement with respect to the claims f) was reached. q was not reached. h N/A. Substance of Interview including description of the general nature of what was agreed to if an agreement was reached, or any other comments: See Continuation Sheet. (A fuller description, if necessary, and a copy of the amendments which the examiner agreed would render the claims allowable, if available, must be attached. Also, where no copy of the amendments that would render the claims allowable is available, a summary thereof must be attached.)

THE FORMAL WRITTEN REPLY TO THE LAST OFFICE ACTION MUST INCLUDE THE SUBSTANCE OF THE INTERVIEW. (See MPEP Section 713.04). If a reply to the last Office action has already been filed, APPLICANT IS GIVEN ONE MONTH FROM THIS INTERVIEW DATE, OR THE MAILING DATE OF THIS INTERVIEW SUMMARY FORM, WICHEVER IS LATER, TO FILE A STATEMENT OF THE SUBSTANCE OF THE INTERVIEW. See Summary of Record of Interview requirements on reverse side or on attached sheet.

PETER PARAS PATENT EXAMINER

Examiner Note: You must sign this form unless it is an Attachment to a signed Office action.

Examiner's signature, if required

Summary of Record of Interview Requirements

Manual of Patent Examining Procedure (MPEP), Section 713.04, Substance of Interview Must be Made of Record

A complete written statement as to the substance of any face-to-face, video conference, or telephone interview with regard to an application must be made of record in the application whether or not an agreement with the examiner was reached at the interview.

Title 37 Code of Federal Regulations (CFR) § 1.133 Interviews

Paragraph (b)

In every instance where reconsideration is requested in view of an interview with an examiner, a complete written statement of the reasons presented at the interview as warranting favorable action must be filed by the applicant. An interview does not remove the necessity for reply to Office action as specified in §§ 1.111, 1.135. (35 U.S.C. 132)

37 CFR §1.2 Business to be transacted in writing.

All business with the Patent or Trademark Office should be transacted in writing. The personal attendance of applicants or their attorneys or agents at the Patent and Trademark Office is unnecessary. The action of the Patent and Trademark Office will be based exclusively on the written record in the Office. No attention will be paid to any alleged oral promise, stipulation, or understanding in relation to which there is disagreement or doubt.

The action of the Patent and Trademark Office cannot be based exclusively on the written record in the Office if that record is itself incomplete through the failure to record the substance of interviews.

It is the responsibility of the applicant or the attorney or agent to make the substance of an interview of record in the application file, unless the examiner indicates he or she will do so. It is the examiner's responsibility to see that such a record is made and to correct material inaccuracies which bear directly on the question of patentability.

Examiners must complete an Interview Summary Form for each interview held where a matter of substance has been discussed during the interview by checking the appropriate boxes and filling in the blanks. Discussions regarding only procedural matters, directed solely to restriction requirements for which interview recordation is otherwise provided for in Section 812.01 of the Manual of Patent Examining Procedure, or pointing out typographical errors or unreadable script in Office actions or the like, are excluded from the interview recordation procedures below. Where the substance of an interview is completely recorded in an Examiners Amendment, no separate Interview Summary Record is required.

The Interview Summary Form shall be given an appropriate Paper No., placed in the right hand portion of the file, and listed on the "Contents" section of the file wrapper. In a personal interview, a duplicate of the Form is given to the applicant (or attorney or agent) at the conclusion of the interview. In the case of a telephone or video-conference interview, the copy is mailed to the applicant's correspondence address either with or prior to the next official communication. If additional correspondence from the examiner is not likely before an allowance or if other circumstances dictate, the Form should be mailed promptly after the interview rather than with the next official communication.

The Form provides for recordation of the following information:

- Application Number (Series Code and Serial Number)
- Name of applicant
- Name of examiner
- Date of interview
- Type of interview (telephonic, video-conference, or personal)
- Name of participant(s) (applicant, attorney or agent, examiner, other PTO personnel, etc.)
- An indication whether or not an exhibit was shown or a demonstration conducted
- An identification of the specific prior art discussed
- An indication whether an agreement was reached and if so, a description of the general nature of the agreement (may be by
 attachment of a copy of amendments or claims agreed as being allowable). Note: Agreement as to allowability is tentative and does
 not restrict further action by the examiner to the contrary.
- The signature of the examiner who conducted the interview (if Form is not an attachment to a signed Office action)

It is desirable that the examiner orally remind the applicant of his or her obligation to record the substance of the interview of each case. It should be noted, however, that the Interview Summary Form will not normally be considered a complete and proper recordation of the interview unless it includes, or is supplemented by the applicant or the examiner to include, all of the applicable items required below concerning the substance of the interview.

A complete and proper recordation of the substance of any interview should include at least the following applicable items:

- 1) A brief description of the nature of any exhibit shown or any demonstration conducted,
- 2) an identification of the claims discussed,
- 3) an identification of the specific prior art discussed,
- 4) an identification of the principal proposed amendments of a substantive nature discussed, unless these are already described on the Interview Summary Form completed by the Examiner,
- 5) a brief identification of the general thrust of the principal arguments presented to the examiner,
 - (The identification of arguments need not be lengthy or elaborate. A verbatim or highly detailed description of the arguments is not required. The identification of the arguments is sufficient if the general nature or thrust of the principal arguments made to the examiner can be understood in the context of the application file. Of course, the applicant may desire to emphasize and fully describe those arguments which he or she feels were or might be persuasive to the examiner.)
- 6) a general indication of any other pertinent matters discussed, and
- 7) if appropriate, the general results or outcome of the interview unless already described in the Interview Summary Form completed by the examiner.

Examiners are expected to carefully review the applicant's record of the substance of an interview. If the record is not complete and accurate, the examiner will give the applicant an extendable one month time period to correct the record.

Examiner to Check for Accuracy

If the claims are allowable for other reasons of record, the examiner should send a litter setting forth the examiner's version of the statement attributed to him or her. If the record is complete and accurate, the examiner should place the indication, "Interview Record OK" on the paper recording the substance of the interview along with the date and the examiner's initials.

Application No. 10/010,065

Continuation of Substance of Interview including description of the general nature of what was agreed to if an agreement was reached, or any other comments: Faxed amendments were not in compliance with Rule 1.121. The amendments faxed 06/10/2003 (which was also mailed), paper #13 and 06/17/2003, paper #14, failed to list the status of claims 1-56. Ms. Verona agreed to send in a new amendment in compliance with Rule 1.21 that reflects the claims as listed in the non-entered amendments (paper # 13 and 14), in light of the last officially entered amendment received 04/14/2003, paper #12. As of 07/08/2003, no amendment has been received and a FINAL office action will be mailed.